Response and Amendment Under §1.116 and

Contingent Suggestion for Declaration of Interference under 37 C.F.R. § 41.202

AMENDMENT TO THE CLAIMS

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of claims

Claims 1-58 (canceled)

- 59. (Previously Presented) A method of inducing apoptosis in mammalian cancer cells comprising exposing mammalian cancer cells to an effective amount of an Apo-2 agonist monoclonal antibody which (a) binds to Apo-2 polypeptide consisting of the contiguous amino acid residues 1 to 411 of SEQ ID NO:1 and (b) induces apoptosis in at least one type of mammalian cancer cell *in vivo* or *ex vivo*.
- 60. (Previously Presented) The method of claim 59 wherein said antibody comprises a singlechain antibody.

Claims 61-64 (canceled)

- 65. (Currently Amended) A method of inducing apoptosis in mammalian cancer cells comprising exposing mammalian cancer cells to an effective amount of an Apo-2 agonist monoclonal antibody which (a) binds to a soluble extracellular domain sequence of an Apo-2 polypeptide consisting of amino acids 54 to 182 of SEQ ID NO:1 and (b) induces apoptosis in at least one type of mammalian cancer cell *in vivo* or *ex vivo*.
- 66. (Previously Presented) A method of inducing apoptosis in mammalian cancer cells comprising exposing mammalian cancer cells to an effective amount of an Apo-2 agonist monoclonal antibody which (a) binds to a soluble extracellular domain sequence of an Apo-2 polypeptide consisting of amino acids 1 to 182 of SEQ ID NO:1 and (b) induces apoptosis in at least one type of mammalian cancer cell *in vivo* or *ex vivo*.

Response and Amendment Under §1.116 and Contingent Suggestion for Declaration of Interference under 37 C.F.R. § 41.202 Claims 67 and 68 (canceled)

- 69. (Previously Presented) The method of claim 59, 65, or 66, wherein said antibody is a chimeric antibody.
- 70. (Previously Presented) The method of claim 59, 65, or 66, wherein said antibody is a humanized antibody.
- 71. (Previously Presented) The method of claim 59, 65, or 66, wherein said antibody is a human antibody.
- 72. (Previously Presented) The method of claim 59, 65, or 66, wherein said antibody comprises an Fab fragment.
- 73. (Previously Presented) The method of claim 59, 65, or 66, wherein said antibody comprises a scFv fragment.
- 74. (Previously Presented) The method of claim 59, 65, or 66, wherein said antibody comprises a F(ab')2 fragment.

Claims 75-78 (canceled)

79. (Previously Presented) The method of claim 59, 65, or 66, wherein said antibody is fused to an epitope tag sequence.

Claims 80-124 (canceled)

125. (Previously Presented) A method of treating cancer comprising exposing mammalian cancer cells to an effective amount of an Apo-2 agonist monoclonal antibody which (a) binds to Apo-2 polypeptide consisting of the contiguous amino acid residues 1 to 411 of SEQ ID NO:1 and (b) induces apoptosis in said mammalian cancer cell *in vivo* or *ex vivo*.

Claim 126 (canceled)

Response and Amendment Under §1.116 and

Contingent Suggestion for Declaration of Interference under 37 C.F.R. § 41.202

- 127. (Previously Presented) The method of claim 125 wherein said agonist antibody is a chimeric antibody.
- 128. (Previously Presented) The method of claim 125 wherein said agonist antibody is a humanized antibody.
- 129. (Previously Presented) The method of claim 125 wherein said agonist antibody is a human antibody.

Claims 130-132 (canceled)

133. (Previously Presented) A method of treating cancer comprising exposing mammalian cancer cells to an effective amount of an Apo-2 agonist monoclonal antibody which (a) binds to a soluble extracellular domain sequence of an Apo-2 polypeptide which consists of amino acid residues 54 to 182 of SEQ ID NO:1 and (b) induces apoptosis in said mammalian cancer cell *in vivo* or *ex vivo*.

Claim 134 (canceled)

- 135. (Previously Presented) The method of claim 133 wherein said agonist antibody is a chimeric antibody.
- 136. (Previously Presented) The method of claim 133 wherein said agonist antibody is a humanized antibody.
- 137. (Previously Presented) The method of claim 133 wherein said agonist antibody is a human antibody.

Claims 138-146 (canceled)

147. (Previously Presented) A method of treating cancer comprising exposing mammalian cancer cells to an effective amount of an Apo-2 agonist monoclonal antibody which (a) binds to a soluble extracellular domain sequence of an Apo-2 polypeptide consisting of

Response and Amendment Under §1.116 and

- Contingent Suggestion for Declaration of Interference under 37 C.F.R. § 41.202 amino acid residues 1 to 182 of SEQ ID NO:1 and (b) induces apoptosis in said mammalian cancer cell *in vivo* or *ex vivo*.
- 148. (Previously Presented) The method of claim 147 wherein said agonist antibody is a chimeric antibody.
- 149. (Previously Presented) The method of claim 147 wherein said agonist antibody is a humanized antibody.
- 150. (Previously Presented) The method of claim 147 wherein said agonist antibody is a human antibody.
- 151. (Previously Presented) The method of claim 125, 133, or 147, wherein said antibody comprises an Fab fragment.
- 152. (Previously Presented) The method of claim 125, 133, or 147, wherein said antibody comprises a scFv fragment.
- 153. (Previously Presented) The method of claim 125, 133, or 147, wherein said antibody comprises a F(ab')2 fragment.
- 154. (Previously Presented) The method of claim 125, 133, or 147, wherein said antibody is fused to an epitope tag sequence.
- 155. (Previously Presented) The method of claim 125, 133, or 147, wherein said mammalian cancer cells are exposed to chemotherapy or radiation therapy.